



Brussels, 18.12.2018
C(2018) 9142 final

COMMISSION IMPLEMENTING DECISION

of 18.12.2018

**amending the marketing authorisation granted by Decision C(2004)3580 for “Alimta -
pemetrexed”, a medicinal product for human use**

(Text with EEA relevance)

(ONLY THE DUTCH TEXT IS AUTHENTIC)

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THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Regulation (EC) No 726/2004 of the European Parliament and of the Council of 31 March 2004 laying down Community procedures for the authorisation and supervision of medicinal products for human and veterinary use and establishing a European Medicines Agency¹, and in particular Article 10 and 28 thereof,

Having regard to Commission Regulation (EC) No 1234/2008 of 24 November 2008 concerning the examination of variations to the terms of marketing authorisations for medicinal products for human use and veterinary medicinal products²,

Having regard to the changes to the terms of the decision granting the marketing authorisation requested by Eli Lilly Nederland B.V. in accordance with Regulation (EC) No 1234/2008,

Having regard to the opinion of the European Medicines Agency, formulated on 18 October 2018 by the Committee for Medicinal Products for Human Use on the periodic safety update report for this medicinal product,

Whereas:

- (1) The placing on the market of the medicinal product "Alimta - pemetrexed" was authorised by Commission Decision C(2004)3580 of 20 September 2004.
- (2) The marketing authorisation holder submitted a periodic safety update report for this medicinal product. This report was assessed by the Pharmacovigilance Risk Assessment Committee as to whether the marketing authorisation concerned should be maintained, varied, suspended or withdrawn.
- (3) The scientific assessment performed by the Committee for Medicinal Products for Human Use, the conclusions of which are set out in Annex IV to this Decision, shows that a decision should be taken amending the marketing authorisation for the medicinal product concerned.

¹ OJ L 136, 30.4.2004, p. 1.

² OJ L 334, 12.12.2008, p. 7.

- (4) The European Medicines Agency's opinion is favourable to the variation of the terms of the decision granting the marketing authorisation as submitted by the marketing authorisation holder.
- (5) Decision C(2004)3580 should therefore be amended accordingly. The Community Register of Medicinal Products should also be updated.
- (6) For the sake of clarity and transparency, it is appropriate, following the amendment of part or parts of the Annexes, to provide for a consolidated version thereof. The Annexes to Decision C(2004)3580 should therefore be replaced.
- (7) The measures provided for in this Decision are in accordance with the opinion of the Standing Committee on Medicinal Products for Human Use,

HAS ADOPTED THIS DECISION:

Article 1

Decision C(2004)3580 is amended as follows:

- 1) Annex I is replaced by the text set out in Annex I to this Decision;
- 2) Annex II is replaced by the text set out in Annex II to this Decision;
- 3) Annex III is replaced by the text set out in Annex III to this Decision.

Article 2

This Decision is addressed to Eli Lilly Nederland B.V., Papendorpseweg 83, 3528 BJ Utrecht, Nederland.

Done at Brussels, 18.12.2018

For the Commission

*Anne BUCHER
Director-General*

